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(71) Applicant (*for all designated States except US*): **CHIESI FARMACEUTICI S.P.A.** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **CAPONETTI, Giovanni** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT). **CATELLANI, Pier, Luigi** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT). **BETTINI, Ruggero** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT). **COLOMBO, Paolo** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT). **VENTURA, Paolo** [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT).

(74) Agents: **MINOJA, Fabrizio et al.**; Bianchetti Bracco Minoja S.r.l., Via Rossini, 8, I-20122 Milano (IT).

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(54) Title: POWDERS CONSISTING OF PARTICLES WITH A PERFECTLY SMOOTH SURFACE, FOR USE AS CARRIERS FOR THE PREPARATION OF INHALATION MIXTURES WITH MICRONIZED DRUGS AND METHOD FOR THEIR PREPARATION

(57) Abstract: Carriers for use in the preparation of mixtures for inhalation powders intended for pulmonary administration of micronized drugs by means of a dry powder inhaler and the method for their preparation are described.

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POWDERS CONSISTING OF PARTICLES WITH A PERFECTLY SMOOTH SURFACE, FOR USE AS CARRIERS FOR THE PREPARATION OF INHALATION MIXTURES WITH MICRONIZED DRUGS AND METHOD FOR THEIR PREPARATION

SUMMARY OF THE INVENTION

The invention relates to carrier particles (the carrier) for use in the preparation of powdery mixtures for inhalation intended for the pulmonary administration of micronized drugs by means of a dry powder inhaler and
5 the method for their preparation.

According to a first aspect, the invention relates to a novel carrier, consisting of a crystalline substance in powder form, in the size range from tens to hundreds of
10 microns, whose particles have a perfectly smooth surface.

A second aspect of the invention relates to a method for smoothing the surface of said particles. The method claimed is able to make the surface of the particles of the carrier smooth, without any roughness, or hollows, clefts and sharp edges, which represent sites of high
15 surface energy to which the drug particles might adhere, without being removed in the aerosol clouds production stage.

The claimed method further permits to improve the uniformity of the surface characteristics of commercially
20 available substances, which are commonly employed as carriers for inhalation powders, whose characteristics are generally rather variable.

Finally, a third aspect of the invention comprises carrier particles that are obtained with the said smoothing method, the surface of which is coated or partially coated with a suitable additive. By virtue of
5 the claimed method the particles of the additive are not released from the carrier particles during inhalation and so do not reach the smaller branching of the pulmonary tree where absorption occurs.

The powders for inhalation obtained by mixing the
10 smooth carrier particles (with or without coating) with a micronized drug give rise to a particularly high respirable fraction of drug.

PRIOR ART

Drugs intended for inhalation therapy, carried out by
15 the administration of dry powders, are characterized by a particle size of a few microns. The particle size is quantified by measuring a characteristic equivalent sphere diameter, known as aerodynamic diameter, which expresses the ability of the particles to be transported
20 as a suspension in an air stream (aerosolization). In general, particles with an aerodynamic diameter of less than 6.4 microns are regarded as respirable, i.e. capable of penetrating into the lungs.

The administration of these drugs in the form of
25 micronized powder requires the use of suitable dry powder inhalers (DPIs).

DPIs in turn can be divided into two basic types:

i) single dose inhalers, for the administration of

single subdivided doses of the active compound;
ii) multidose dry powder inhalers (MDPIs), pre-loaded with quantities of active principles sufficient for longer treatment cycles.

5 Although micronization of the drug particles is essential for penetration to the deepest branchings of the pulmonary tree during inhalation, it is also known that the finer are the particles, the stronger are the cohesion forces. In multidose inhalers, said effects
10 hamper the loading of the doses of powder from the reservoir system to the aerosolization chamber, since the cohesion forces reduce free flowing of the particles and promote their agglomeration and/or their adhesion to the walls. The aforementioned effects therefore impair the
15 efficiency and reproducibility of the delivered dose and are detrimental to the respirable fraction.

Multidose inhalers work properly when so-called free-flowing powders are used, generally formulated by mixing the micronised drug with a carrier material (generally
20 lactose, preferably α -lactose monohydrate) consisting of coarser particles, approximately equal or greater than 100 microns. In such mixtures, the micronised active particles mainly adhere to the surface of the carrier particles whilst in the inhaler device; on the contrary,
25 during inhalation, a redispersion of the drug particles from the surface of the carrier particles occurs allowing the formers to reach the absorption site into the lungs.

Mixing with the carrier also facilitate the

introduction and withdrawal of the inhalation preparation, in a regular dose, from the reservoir of a multidose inhaler or its dosage in single-dose containers. Mixing of the micronized drug with the coarse carrier therefore leads to the production of a mixture in which the micronized drug is distributed uniformly on the carrier particles as a result of the interactions, usually of an electrostatic nature, which establish between the drug particles and the carrier particles. Said interactions lead to the production of a so-called ordered mixture. It is extremely important for the interactions to be weak and reversible, so that, since transport in the air stream and the respirability of the powder depend on the particle size, only the micronized drug particles will be able to be deposited in the lungs, whereas the coarser carrier particles will be deposited, because of their mass, in the upper airways. Due to the weak interactions between the two components of the mixture, breathing-in through the inhaler causes separation of the micronized drug particles from the coarse carrier particles and therefore inhalation of the smaller particles and deposition of the coarser particles in the oropharyngeal cavity. Accordingly, it is of great applicative interest to find new carriers for inhalers and new techniques for the production of drug-carrier mixtures easy to handle and able to generate a high respirable fraction.

The use of a carrier is indeed not free of drawbacks

in that the strong interparticle forces between the two ingredients may prevent the separation of the micronised drug particles from the surface of the coarse carriers ones on inhalation, so compromising the availability of the drug to the respiratory tract.

In the prior art there are many examples of processes for modifying the surface conditions of the carrier with the aim of reducing the strength of the interactions between the particles during inhalation, without causing pre-separation of the drug particles in the inhaler.

Ganderton (GB 2 240 337) reports that the surface conditions of the particles, in particular their rugosity, are critical for the behaviour of the carrier during inhalation and claims pharmaceutical carriers, such as lactose, consisting of particles whose rugosity is controlled by a cristallization process. The rugosity of the said particles is evaluated using measurements of surface area, based on gas permeametry. The surface area value measured by this technique, relative to the theoretical surface area value, provides a numerical index of rugosity called Ganderton scale.

It is known anyway that measurements based on the said parameter (gas permeametry) only provide reliable data in the case of powders consisting of particles with diameter below 45 μm (subsieve range). In fact, by using such method, no difference between the lactose before and after the smoothing treatment can be detected in the case of particles with a mean diameter of about 100 μm .

Moreover the values obtained are not reliable (≈ 2.5) as demonstrated in Example 1.

In any case, the method of the prior art makes it possible only to reduce the surface rugosity of the carrier particles, as they can have a degree of surface rugosity up to 1.75 , determined according to the permeametry method.

Staniforth (WO 95/11666) claims a milling process preferably carried out in a ball mill, called corrasion (for analogy with the effect of wind on rocks), which alters the surface characteristics of the carrier by removing asperities in the form of small grains; these grains in turn can become attached to the clefts of the surface area of the particles, so saturating the high-energy sites. As a result of this preliminary treatment of the carrier, the micronized drug particles are deposited preferentially on lower-energy sites and so are subject to weaker forces of interparticle adhesion.

It is also known from the literature that various types of commercial lactose can have a moderate degree of surface rugosity.

In Kawashima et al. (Int J Pharm 172, 1998, 179-188) examples are given of crystalline lactose with rugosity between 1.33 and 1.13, evaluated on the basis of the perimeter of the particles determined by scanning electron microscope (scale utilized by Kawashima).

Podczeck F (J Adhesion Sci Technol 12, 1998, 1323-1339) reports that Pharmatose 125 M (a commercially

available lactose) is characterized by a surface rugosity, expressed in μm , of 1.12 ± 0.74 (scale utilized by Podczeck).

5 The values reported relate however to batches of lactose with a granulometric distribution between approx. 30 and 90 μm and characterized by a median diameter of approx. 60 μm . It is known, however, that the finer the particles, the more they have a regular shape and so are intrinsically characterized by a lower rugosity value.

10 On the other hand, the operation of some multidose inhalers requires the use of optimum carriers of high flowability, a characteristic that can only be imparted by using particles with a greater granulometric distribution.

15 Disaggregation of the active principle from the carrier during inhalation can also be made more efficient by addition of a fraction of fine particles of the same carrier. The Boheringer patent EP 0 663 815 claims the use of carriers for controlling and optimizing the amount
20 of drug released during the aerosolization phase, comprising suitable mixtures of coarse particles with size $> 20 \mu\text{m}$ and of fine particles with size $< 10 \mu\text{m}$.

Finally, in the prior art, additives with lubricant, glidant or anti-adherent properties, dry-mixed with the
25 carrier, have been employed with the aim of reducing the forces of attraction between drug and carrier. For example, mixing of magnesium stearate with crystalline lactose is able to reduce the forces of adhesion between

drug and carrier, when this mixture is used as inhalation carrier. For explaining the effectiveness of magnesium stearate in the aerosolization of inhalation powders, investigations conducted on powder mixtures for tablets
5 can be taken into account (Staniforth et al., J. Pharm. Pharmacol. 1982, 34, 141-145). These investigations showed that the presence of lubricants causes a decrease in cohesion of the tablets because they form a lubricated layer on the powder particles that are to be pressed
10 together, thereby interfering with the bond between them. This mechanism is also regarded as responsible for the decrease in strength of adhesion of the micronized drug particles on the carrier particles (Kassem, thesis, London University, 1990).

15 In WO 96/23485, the particles are mixed with a substance with anti-adherent or antifriction properties, consisting of one or more compounds selected from amino acids (in particular leucine), phospholipids or surfactants; deposition of the additive on the carrier is
20 preferably carried out in the dry form, and does not give rise to a complete coating of the carrier, but rather to a discontinuous covering in order to saturate the high-energy sites. Preferably, the carrier particles and the additive are submitted to the corrosion process in a ball
25 mill as described in WO 95/11666.

It follows from examination of the prior art that in the case of an inhalation powder, consisting of a drug-carrier mixture, efficient disaggregation of the active

principle from the carrier during inhalation is dependent upon the drug-carrier interparticle forces and so depends on the surface characteristics of the latter.

Furthermore, it has been found in certain cases that commercial batches of lactose obtained from the same manufacturer, though possessing the same physicochemical and technological characteristics, exhibited substantially different behaviours on inhalation, so that they could not be regarded as equivalent. A difference in surface area among these batches can be for instance detected by the multiple-point BET method, even in cases when they could not be appreciated by gas permeametry or by the single-point BET method.

Images obtained with the scanning electron microscope showed, in turn, that this difference was to be ascribed to the different conditions of surface rugosity of the particles.

In the batches of lactose examined, it was also noted that there was a different percentage and granulometric distribution of fine particles.

The presence of fine particles in the lactose for inhalation might be useful for optimizing the respirability of an active principle mixed with a coarse carrier, as claimed in patent EP 0 663 815. However, since only the fine fraction below 10 μm is effectively responsible for the decrease in the interparticle forces, whereas the fraction greater than 10 μm lowers the flowability of the powder, it is important to be able to

control the percentage and distribution of the fine particles in accordance to the use the carrier is directed to.

As already observed, the commercially available
5 excipients, being substances widely used in the pharmaceutical field and intended for several applications, exhibit small but substantial variations, e.g. of surface area or distribution of fine particles, which can impair their performance when they are used for
10 particular purposes, such as carriers for inhalation powders.

Although it has been widely reported that by altering the surface characteristics of the carrier it is possible to increase the respirable fraction of the inhaled drug,
15 it has never been previously described a process of treatment of carrier particles for inhalation powders able to eliminate the random variations of the surface characteristics caused by the ordinary manufacturing processes.

20 OBJECT OF THE INVENTION

It has now been found that by pre-treating the carrier in a high-speed mixer-granulator it is possible to obtain carrier particles with median diameter greater than 90 μm , smooth and homogeneous, characterized by a
25 surface rugosity of less than 1.1, and preferably between 1.0 and 1.1 (fractal scale), suitable for making inhalation powders.

A first aspect of the invention therefore relates to

a new type of carrier for inhalation powders, characterized by particles having perfectly smooth surfaces and rounded edges or corners. In these particles, surface rugosity - which often creates problems when using the carriers for the preparation of inhalation powders - has been completely eliminated, making the particles perfectly smooth.

A second aspect of the invention relates to the method of preparation of these carriers consisting of particles with perfectly smooth surface. The claimed method allows to obtain said smooth particles starting from an industrial powder consisting of rough particles, without substantially altering their average size and their geometry.

The carrier of the invention can be prepared using a high-speed mixer-granulator, an apparatus designed and normally used for agglomerating solid particles and not for smoothing them individually. This generally consists of a cylindrical chamber with a chamfered bottom, in which a rotating paddle is inserted, and once this is running at a suitable speed it causes the powder contained within the cylindrical chamber to roll along the chamber walls. The mixing chamber is sealed by a cover, which contains a sprayer for adding liquid, and can operate in controlled conditions of temperature and pressure. Until now, this type of equipment has been used exclusively for the preparation of granules or pellets, i.e. to agglomerate the individual particles, by means of

a liquid binder, to give more complex structures, called granules.

It has now been found that in certain conditions, the use of such apparatus allows to alter the surface characteristics and shape of particles of pharmaceutical excipients, such as those proposed as carriers for inhalation powders, without agglomerating them and without significantly changing their crystalline structure and physicochemical properties. The process of the invention, which has been called "particle smoothing", makes it possible to alter the surface characteristics of the carrier particles for inhalation use, without causing their agglomeration into granules or pellets, and is explained in detail in the examples given below. As a result of this smoothing process, a powder is obtained that consists of individual particles. Moreover, the process only gives rise to a slight reduction of the particle size relevant to the starting product, without increasing the fraction of fine particles. The carrier of the invention, consisting of individual smooth particles, is therefore clearly different from carriers in which the particles are agglomerated to form pellets, i.e. granular shapes obtained by agglomerating several individual particles together, as those described in WO 95/24889, WO 95/09615, WO 95/09616 .

The method of the invention is rapid and convenient, and produces particles with smooth surfaces with rounded edges and corners, eliminating from their surface the

asperities and clefts originally present, i.e. high-energy sites on which the active particles can strongly adhere. Furthermore, as a result of the claimed process that consists of stages of wetting and slow drying, as will be explained in more detail later, the fine particles present in the powder prior to be smoothed are eliminated from the final product. The method of the invention therefore makes it possible to eliminate the small random differences that are inherent to the industrial manufacturing process, and are detrimental to their efficiency when they are used as carriers for inhalation powders.

A further advantage of the method of the invention is that it makes possible to obtain particles that are perfectly smooth and are therefore characterized by a very low surface rugosity, starting from particles with a coarser particle size, for instance between 30 and 600 μm (preferably between 90 and 150 μm).

The smoothing process is carried out in repeated stages of wetting with a solvent and drying of the carrier particles. For wetting the particles an aqueous or aqueous-alcoholic solution is preferably used, and the powder particles are caused to roll on the walls of the mixing chamber as it is submitted to rapid mixing by the rotating paddle of the high-speed granulator. Photographs taken with the scanning electron microscope (SEM), of samples of lactose particles taken at the beginning (a), during (b and c) and at the end (d) of the smoothing process, are shown

(Fig. 1). This method leads to surface smoothing of the lactose particles, eliminating the sharp edges and any fissures, without significantly changing the size and shape of the particles. The method also leads to
5 elimination of the fraction of fine particles that were present in the original powder. The graphs in Fig. 1, superimposed on the images of the particles, provide a quantitative evaluation of the change in rugosity as a result of the smoothing process, by means of an image
10 analysis system applied to the photographs of the particles. This approach makes it possible to measure, on the surface of the particles, the variation in level of grey as a function of position (the region measured for each particle is indicated by the shaded rectangle
15 reproduced in the Figure). The surface rugosity of the particles is quantified from the deviation from linearity of the lines reproduced in the diagram, which relate to the variation in grey level of the image relating to the surface of the particle. The smoother is the particle,
20 the more uniform is the level of grey of its image, and hence more straight is the line. Quantitative measurement of rugosity can be obtained from measurement of the fractal dimension of this line which expresses the level of grey as a function of the position on the surface of
25 the crystal (fractal scale). The more this line moves away from a straight line, the more the value of the fractal dimension is different from that of the topological dimension, which is the unity for a straight

line. The fractal dimension of the smooth particles is indeed equal to one, as explained by B.H. Kaye (Part. Part. Syst. Charact. 1993, 10, 99-110).

5

Table 1

Stages of processing	Fractal dimension
Starting lactose	1.23
Step # 1	1.08
Step # 7	1.06
Smoothed lactose	1

10

Table 1 shows the values of fractal dimension corresponding to the "rugosity" lines shown in Fig. 1, according to the various stages of smoothing indicated. The particles are considered to be acceptably smooth when the fractal dimension is ≤ 1.1 .

15

20

Surface smoothing, redistribution of the fraction of fine particles in the fissures and rounding of the edges are obtained by a mechanism of controlled solubilization and deposition of the lactose, due to wetting with the solvent sprayed on the powder followed by slow drying, while the mixture is submitted to rapid mixing. Since the chamber of the mixer is of cylindrical shape and the rotating paddle (impeller) is positioned centrally at the base of the chamber, the powder is caused to roll on the walls of the said chamber. During this rolling, rounding

of the edges of the individual particles occurs, without causing any agglomeration of the particles on account of the choice of type and amount of solvent used. The surface characteristics of the carrier particles are a direct consequence of the method of treatment. The final result is the reduction or elimination of the high-energy sites from the surface of the particles, which arise in the original operations of manufacturing of the carrier. Smoothing the lactose particles by means of the method of the invention makes it possible to obtain a carrier for inhalation powders of higher performances with respect to the respirable fraction of drug, in comparison with powder that has not been smoothed.

Another aspect of the present invention relates to the preparation of smooth powders for inhalation purposes using a solution or suspension of the carrier in a liquid containing a suitable additive with properties such as to promote disaggregation of the drug from the surface of the carrier during inhalation. With respect to the prior art, the process of the invention permits to achieve an at least partial coating of the carrier particles in liquid phase. Advantageously during the smoothing process, the particles of additive are trapped on the surface of the carrier particles, forming a coating capable of modulating the drug-carrier interparticle forces. By virtue of said process, the particles of additive are firmly bound to the surface of the carrier. As a consequence, during inhalation, the particles of

additive are deposited together with the coarse carrier particles in the oropharyngeal cavity, so they are not inhaled.

DESCRIPTION OF THE PREFERRED CONDITIONS

5 The carrier particles can be made of any physiologically acceptable, inert material. Preferred carriers are those consisting of crystalline sugars such as glucose, mannose, galactose, sorbitol, mannitol, lactose, saccharose, trehalose, raffinose, cyclodextrins
10 and mixtures thereof. Carriers comprising α -lactose monohydrate are even more preferred. The particles of the carrier can have a size between 30 and 600 μm , preferably between 90 and 150 μm .

 To carry out the process of the invention, high-
15 speed mixer-granulators such as the Roto J Zanchetta (Zanchetta S.p.A. Lucca), or the Diosna P50 (Dierksöhne, Osnabruck, Germany) can be advantageously used.

 Short-chain aliphatic alcohols, or water-alcohol mixtures, can be used advantageously as solvents. The
20 preferred conditions envisage the use of a water:alcohol mixture from 9:3 to 3:4 v/v; more preferred conditions are those in which the water:ethanol ratio is 5:3 v/v; even more preferred are those in which the ratio is 3:3 v/v.

25 The optimum ratio between volume of solution and amount of carrier to be smoothed is between 5 ml/100 g and 70 ml/100 g, preferably 53 ml/100 g.

 The operating conditions of the high-speed mixer-

granulator are: rotary speed of the mixing paddle (impeller) between 25 and 600 rev/min; temperature of the outer jacket of the mixing chamber between 20°C and 90°C, preferably 50°C; solution nebulization pressure between 1 and 10 bar, preferably 3.5 bar; diameter of the nebulization nozzle between 0.7 and 2.0 mm, preferably 1.0 mm; vacuum inside the mixing chamber between -0.2 and -0.8 bar, preferably -0.7 bar.

The total mixing time of the carrier powder particles during smoothing is preferably comprised between 120 and 300 minutes, with a period of drying, after each stage of spraying of the smoothing solution, of between 10 and 30 minutes.

The additives can be selected from those belonging to the class of lubricants, such as magnesium stearate, sodium benzoate, sodium stearyl fumarate or to the class of anti-adherents such as leucine and isoleucine. Other additives that can be advantageously used are soluble polymers such as hydroxyethylcellulose, methylcellulose, carboxymethylcellulose, polyvinylpyrrolidone, polyethyleneglycol, cyclodextrins.

With respect to the volume of the solution, the additive is added in amounts preferably between 0.05% and 2%, more preferably between 0.25% and 1%, even more preferably 0.45%

The final formulation should contain from 0.1 to 1% by weight of a lubricant, more preferably 0.25% by weight of magnesium stearate.

The spraying times of the solution or suspension of additive are between 5 and 30 seconds.

The inhalation formulations that can be obtained by mixing active principles in the form of micronized powder
5 (median diameter $\leq 6.4 \mu\text{m}$) with carrier powders obtained by the claimed method constitute a further object of the invention.

Preferred active ingredients are the drugs usually administered in micronized form for the treatment of
10 respiratory diseases by inhalation, for example steroids such as beclometasone dipropionate, flunisolide and budesonide, β -agonists such as salbutamol, formoterol, salmeterol, terbutaline, their salts and mixtures thereof, anticholinergics such as ipratropium bromide and
15 oxitropium bromide. However, any active principle suitable for endobronchial administration can be used.

The process of the invention is illustrated in detail by the examples that follow, which do not limit its application in any way.

20 Fig. 1 and Table 1 give the fractal dimensions relating to particle surface rugosity of α -lactose monohydrate before, during and at the end of smoothing.

Example 1: Inhalation powder made of beclometasone dipropionate (BDP) mixed with smoothed α -lactose monohydrate carrier.
25

Method of manufacture

a) The carrier particles of the invention were prepared as follows:

750 g of lactose α -monohydrate (size fraction 90-150 μm) was loaded into the high-speed granulator (Zanchetta Roto Junior, Lucca, I), with the heating jacket at a temperature of 50°C and with vacuum of -0.7 bar. After starting the mixing impeller at a speed of 50 rev/min, 40 ml of a 3:5 v/v mixture of ethyl alcohol:distilled water was sprayed on the powder for a time of 10 seconds. After completion of the spraying stage, the speed of the mixing impeller was increased to 450 rev/min for 15 minutes, in order to cause the powder particles to roll along the circular wall of the mixer and promote drying of the said powder particles. The stages of spraying and drying, as described, were repeated 10 times, until a total solution volume of 400 ml was added.

b) At the end of the smoothing stage, the powder was removed from the mixer chamber and was placed in an air-circulation stove for 240 minutes at 60°C.

c) Afterwards, the dried powder particles were placed and kept in a climate chamber at a temperature of 20°C and 60% RH. The smoothed lactose particles, photographed using a scanning electron microscope, were found to possess a morphology with rounded corners and a smooth surface, as can be seen in Fig. 1, which shows the variation in rugosity of the lactose particles during the smoothing stages.

The size of the lactose particles before and after smoothing, expressed as geometric mean diameter (d_g) and geometric standard deviation (σ_g) had not been altered

substantially:

starting lactose d_g : 123.9 μm ; σ_g : 1.05

lactose after smoothing d_g : 119.3 μm ; σ_g : 1.10

The rugosity factor calculated by SEM is reported in
5 Table 1.

The rugosity factor was also calculated according to
Ganderton (GB 2 240 237), i.e. by calculating the ratio
between the specific surface area, determined by air
permeametry and the geometric specific surface area.

10 The results are reported in Table 2.

Table 2

Composition	Rugosity Factor
BDP/lactose	2.7
BDP/smoothed lactose	2.4

The results demonstrate that using the specific surface
15 area obtained by the air permeametry method described in
GB 2 240 237, no difference between the original and the
smoothed lactose can be detected. Moreover, in both
cases, using as a starting carrier particles of size
greater than 90 μm , the rugosity factor turned out to be
20 greater than 1.75, which is the upper limit value of the
aforementioned patent.

In order to check for potential change in the crystalline
structure, powder X-ray diffraction and differential
scanning calorimetry analysis were carried out on the

starting lactose and on the smoothed lactose. Both the thermal and diffraction patterns demonstrated that the crystalline structure of the lactose had not been changed by the process of smoothing.

- 5 d) The inhalation powder, consisting of a drug/carrier mixture, was prepared as follows:

0.684 g of BDP was mixed with 80 g of smoothed lactose, in three different steps:

- 10 i. pre-mixing for 10 minutes of the whole amount of BDP with approx. 9/10 of the carrier, in a rotating-chamber mixer (Turbula®), using a cylindrical steel container with volume of approx. 300 ml;
- ii. disaggregation of the aggregates on a metal screen with 300 µm holes;
- 15 iii. "washing" of the screen used for disaggregation with the remainder of the carrier and mixing of all the combined powder for 30 minutes in the steel container of the Turbula® mixer.

20 The dose of inhalation mixture that was loaded in the inhaler for delivery was made up of 24.8 mg of carrier and 0.2 mg of BDP.

Monitoring of the respirable fraction of the inhalation powder

25 A quantity of inhalation powder was loaded in a Pulvinal® powder inhaler (Chiesi Farmaceutici, Italy) and kept in a climate chamber at 20°C and 60% RH for about 3 days before starting each aerosolization test.

The respirable fraction of the inhalation powder

prepared according to Example 1 was evaluated using a twin-stage impinger (Apparatus of type A for the aerodynamic evaluation of fine particles described by the FU IX, 4th supplement 1996). The instrument is made of various glass components, connected together to form two chambers capable of separating the inhalation powder as a function of its aerodynamic dimensions, called the upper separating chamber (Stage 1) and the lower separating chamber (Stage 2). A rubber adapter provides connection to the Pulvinal® inhaler that contains the dose of inhalation powder. The apparatus is connected to a vacuum pump that produces a flow of air through the separating chambers and the inhaler connected to them. During the delivery stage, the differences in shape and size of the particles of the inhalation powder, as well as the air stream, cause discrete deposition of the particles in the separating chambers. Inserting the inhaler in the appropriate rubber adapter and applying an air flow of 60 l/min for 10 seconds (a flow that reproduces the inspiratory capacity of an asthmatic patient), it was possible to separate the particles of the powder delivered in the two stages of the apparatus (Stage 1 and Stage 2). The limit value of aerodynamic diameter, d_{ae} , for deposition in the lower separating chamber is 6.4 μm . Particles with larger d_{ae} are deposited in Stage 1 and particles with smaller d_{ae} in Stage 2. A minimum volume of solvent is present in both stages (30 ml in Stage 2 and 7 ml in Stage 1), consisting of an acetonitrile:water

mixture 6:4 v/v, to prevent impaction of the particles on the apparatus walls and to facilitate their recovery.

After nebulization of a dose of powder in the twin-stage impinger, the apparatus is dismantled and the quantities of drug deposited in the two separating chambers are recovered by washing with solvent mixture and brought up to a volume of 100 ml in two graduated flasks, one for Stage 1 and one for Stage 2, respectively. The quantities collected in the two flasks are then analysed.

Results

The results relating to determination of the respirable fraction are given in Table 3 in comparison with a similar preparation obtained by mixing BDP and unsmoothed lactose (reference preparation). The respirable fraction is expressed as the percentage of particles of active principle found in Stage 2 relative to the quantity of active principle delivered.

The results demonstrate a significant increase in respirable fraction.

Table 3

Composition	Respirable fraction (%)
BDP/lactose	9.0 ± 1.0
BDP/smoothed lactose	21.8 ± 2.7

Example 2: Inhalation powder of beclometasone dipropionate mixed with smoothed α -lactose monohydrate carrier in the presence of isoleucine.

5 a) The carrier particles of smoothed lactose in the presence of isoleucine were prepared as described in Example 1 (a), coating the particles using an aqueous-alcoholic solution (5:3 v/v) containing isoleucine equal to 0.75% of the amount of lactose used. Fig. 2 shows the smoothed particles and graphic evaluation of their
10 rugosity.

b) The inhalation powder was prepared by mixing BDP with smoothed lactose in the presence of isoleucine according to the method described in Example 1 (d).

15 c) Determination of the respirable fraction of the prepared mixture was effected using the Pulvinal® powder inhaler, in accordance with the method described in Example 1.

The results are given in Table 4.

20 The results show that there is a significant increase in respirable fraction, as well as with respect to the preparation containing smoothed lactose with no additive (Table 3).

Table 4

Composition	Respirable fraction (%)
BDP/lactose	9.0 \pm 1.0
BDP/smoothed lactose with isoleucine	30.5 \pm 2.8

Example 3: Inhalation powder of beclometasone dipropionate mixed with smoothed α -lactose monohydrate carrier in the presence of magnesium stearate.

- 5 a) The carrier particles of smoothed lactose in the presence of magnesium stearate were prepared as described in Example 1 (a), coating the particles using an aqueous-alcoholic suspension (5:3 v/v) containing magnesium stearate equal to 0.25% of the amount of lactose used. Fig. 3 shows the smoothed particles and graphical
10 evaluation of their rugosity. The microscope images also demonstrate absence of fine particles adhering to the surface of the smoothed crystal of lactose, indicating that the particles of magnesium stearate remain trapped on the smoothed surface.
- 15 b) The inhalation powder was prepared by mixing BDP with smoothed lactose in the presence of magnesium stearate according to the method described in Example 1 (d).
- c) Determination of the respirable fraction was effected using the Pulvinal[®] powder inhaler, in accordance with
20 the method described in Example 1.

The results are presented in Table 5. Also in this case, a significant increase in respirable fraction was observed, as well as with respect to the preparation containing smoothed lactose with no additive.

Table 5

Composition	Respirable fraction (%)
BDP/lactose	9.0 ± 1.2
BDP/smoothed lactose with magnesium stearate	31.0 ± 1.8

Example 4: Inhalation powder of beclometasone dipropionate mixed with smoothed α -lactose monohydrate carrier in the presence of magnesium stearate.
5 Performances at different air flows.

a) The carrier particles of the invention were prepared as follows:

- a) 750 g of lactose α -monohydrate (size fraction 90-150 μ m)
10 was loaded into the high-speed granulator (Zanchetta Roto Junior, Lucca, I), with the heating jacket at a temperature of 50°C and with vacuum of -0.7 bar. After starting the mixing impeller at a speed of 50 rev/min, 40 ml of a 1:1 v/v mixture of ethyl alcohol:distilled water,
15 containing magnesium stearate equal to 0.25% of the amount of lactose used, was sprayed on the powder for a time of 10 seconds. After completion of the spraying stage, the speed of the mixing impeller was increased to 450 rev/min for 15 minutes, in order to cause the powder
20 particles to roll along the circular wall of the mixer and promote drying of the said powder particles. The stages of spraying and drying, as described, were

repeated 10 times, until a total solution volume of 400 ml was added.

b) At the end of the smoothing stage, the powder particles were removed from the mixer chamber and was placed in an air-circulation stove for 240 minutes at 60°C.

c) Afterwards, the dried powder was placed and kept in a climate chamber at a temperature of 20°C and 60% RH.

d) The inhalation powder was prepared by mixing BDP with smoothed lactose in the presence of magnesium stearate according to the method described in Example 1 (d).

e) Determination of the respirable fraction of the prepared mixture was carried out using the Pulvinal® powder inhaler, in accordance with the method described in Example 1. The respirable fraction was also determined after applying an air flow of 30 l/min.

The results are given in Table 6, in comparison with the reference preparation. The results show that there is a significant increase in respirable fraction at both air flows. This allows the powder of the invention to be used with medium-resistance as well as high-resistance dry powder inhalers.

Table 6

Composition	Respirable fraction (%) (30 L/min)	Respirable fraction (%) (60 L/min)
BDP/lactose	—	9.0 ± 1.0
BDP/smoothed lactose with magnesium stearate	62.7 ± 1.0	68.6 ± 1.5

Example 5: Inhalation powder of low beclometasone dipropionate dose mixed with smoothed α -lactose monohydrate carrier in the presence of magnesium stearate.

a) The carrier particles of smoothed lactose in the presence of magnesium stearate were prepared as described in Example 4 (a).

b) 0.068 g of BDP was mixed with 80 g of smoothed lactose in presence of magnesium stearate, in three separate steps:

i. pre-mixing for 10 minutes of the whole amount of BDP with approx. 9/10 of the carrier, in a rotating-chamber mixer (Turbula®), using a cylindrical steel container with volume of approx. 300 ml;

ii. disaggregation of the aggregates on a metal screen with 300 μ m holes;

iii. "washing" of the screen used for disaggregation with the remainder of the carrier and mixing of all the combined powder for 60 minutes in the steel container of the Turbula® mixer.

The dose of inhalation mixture that was loaded in the

inhaler for delivery was made up of 24.98 mg of carrier and 0.02 mg of BDP.

c) Determination of the respirable fraction of the prepared mixture was effected using the Pulvinal[®] powder inhaler, in accordance with the method described in Example 1.

The results are given in Table 7, in comparison with the reference preparation. The results show that there is a significant increase in respirable fraction and therefore the process of the invention is also suitable for preparing powders for inhalation containing low-strength active ingredients.

Table 7

Composition	Respirable fraction (%)
BDP low dose/smoothed lactose with magnesium stearate	46 ± 12.4

CLAIMS

1. Particles for use as a carrier in the preparation of pharmaceutical formulations for the pulmonary administration of micronized drugs by means of a powder inhaler, wherein the median diameter is greater than 90 μm and the surface rugosity is less or equal to 1.1.
2. Particles according to Claim 1, consisting of one or more saccharides selected from glucose, mannose, galactose, sorbitol, mannitol, lactose, saccharose, trehalose, raffinose and cyclodextrins.
3. Particles according to Claims 1 and 2, consisting of α -lactose monohydrate.
4. Particles according to the preceding claims with a starting diameter between 30 and 600 μm .
5. Particles for use as a carrier according to claim 1, wherein the surface is coated with an additive selected from lubricants, anti-adherents and soluble polymers.
6. Particles according to Claim 5 in which the lubricant is magnesium stearate, sodium benzoate, or sodium stearyl fumarate.
7. Particles according to Claim 5 in which the anti-adherent is leucine or isoleucine.
8. Particles according to Claim 5 in which the soluble polymer is hydroxyethylcellulose, methylcellulose, carboxymethylcellulose, polyvinylpyrrolidone, polyethyleneglycol, or cyclodextrins.

9. A method of preparation of smooth carrier particles wherein smoothing of the particles is accomplished after repeated stages of wetting with a solvent and drying.

10. A method according to Claim 9, carried out using a high-speed granulator consisting of a cylindrical mixing chamber (container) in which a rotating paddle (impeller) and a spray nozzle are inserted, capable of operating in controlled conditions of temperature and pressure.

11. A method according to Claims 9 and 10, wherein the mixing time is between 120 and 300 minutes.

12. A method according to Claims 9-11 wherein a carrier substantially free from fine particles is obtained.

13. A method according to Claims 9-12, wherein the solvent used for smoothing contains, dissolved or dispersed, an additive in an amount between 0.05 and 2%.

14. A method for the preparation of pharmaceutical formulations wherein one or more active ingredients whose particles have a median diameter $\leq 6.4 \mu\text{m}$ are added to the carrier, prepared according to any one of the preceding claims.

15. A method according to Claim 14, wherein the active ingredient is a β -agonist.

16. A method according to Claim 15, wherein the active ingredient is salbutamol, formoterol, salmeterol, terbutaline, their salts and their epimers.

17. A method according to Claim 14, wherein the active ingredient is an anti-inflammatory steroid.

18. A method according to Claim 17, wherein the active

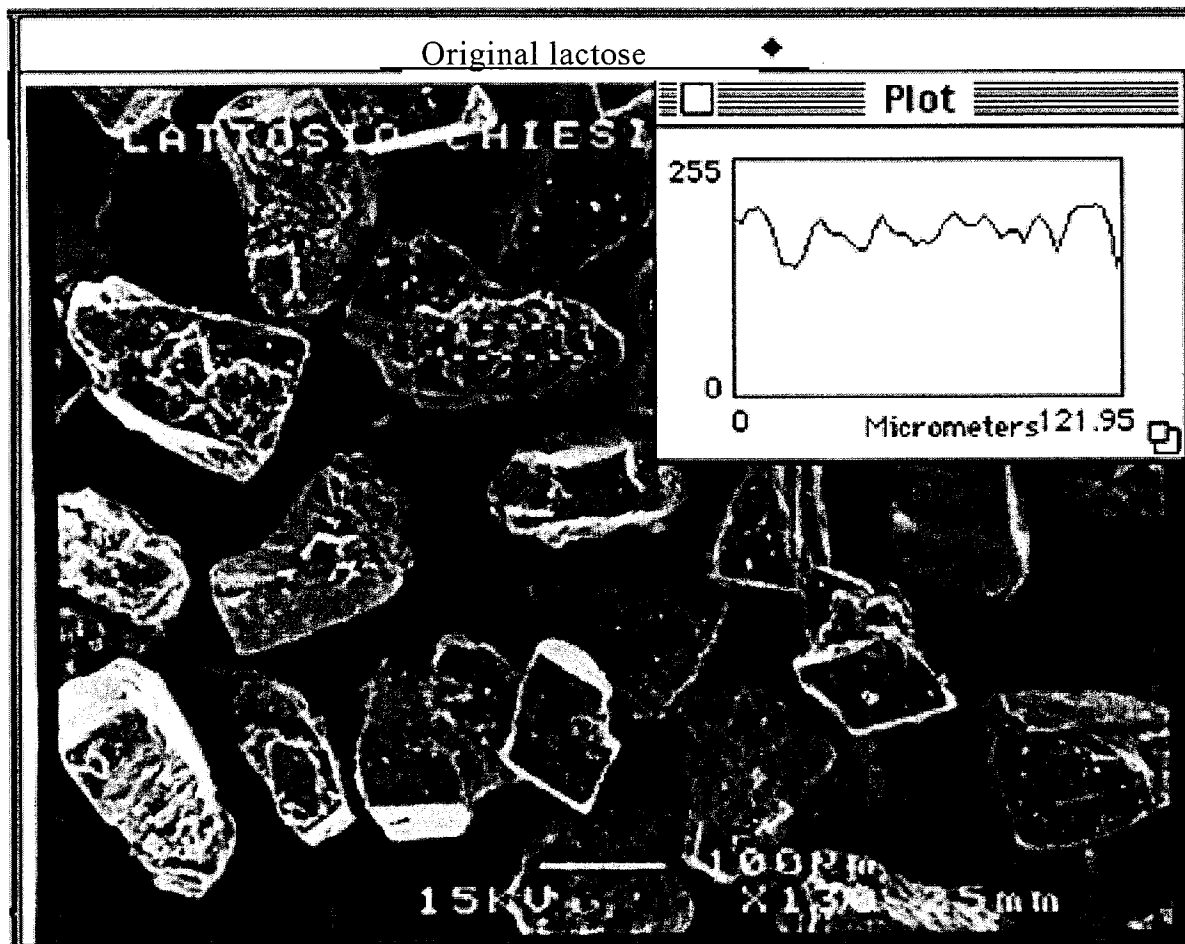
ingredient is beclometasone dipropionate, flunisolide, budesonide and their epimers.

19. A method according to Claim 14, wherein the active ingredient is an anticholinergic.

5 20. A method according to Claim 19, wherein the active ingredient is ipratropium bromide or oxitropium bromide.

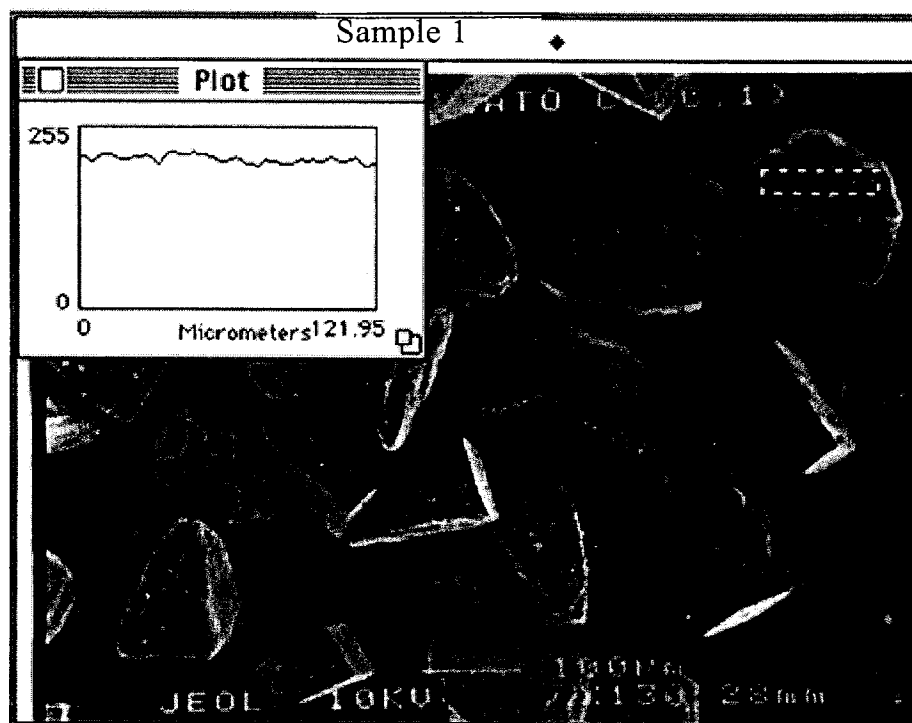
21. Pharmaceutical compositions for inhalation, obtained by mixing active principles in the form of micronized powder with the particles of Claims 1-8.

FIGURE 1

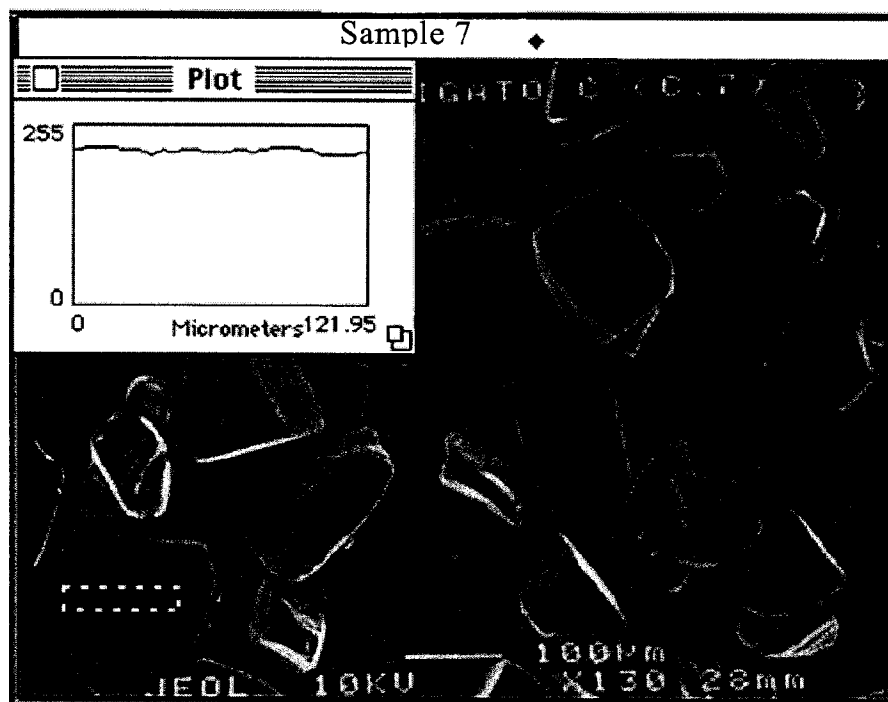


a

FIGURE 1

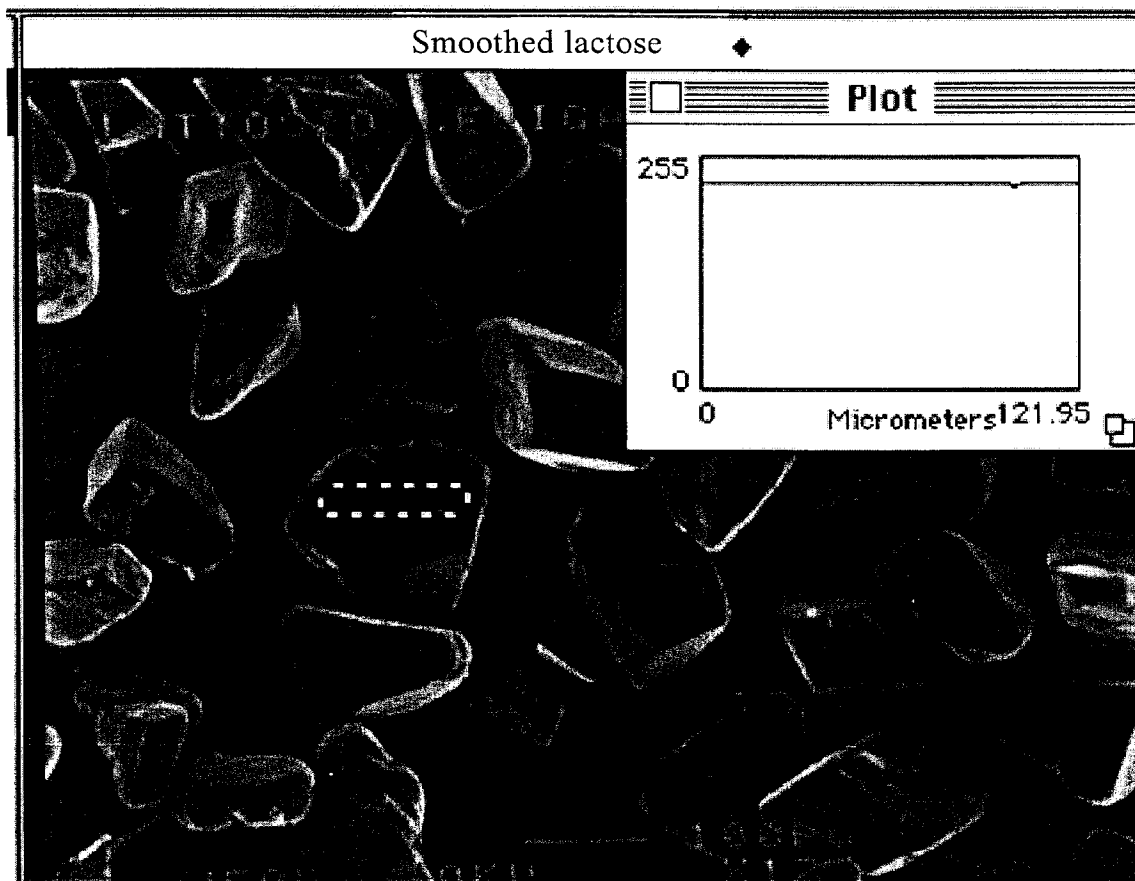


b



c

FIGURE 1



d

FIGURE 2

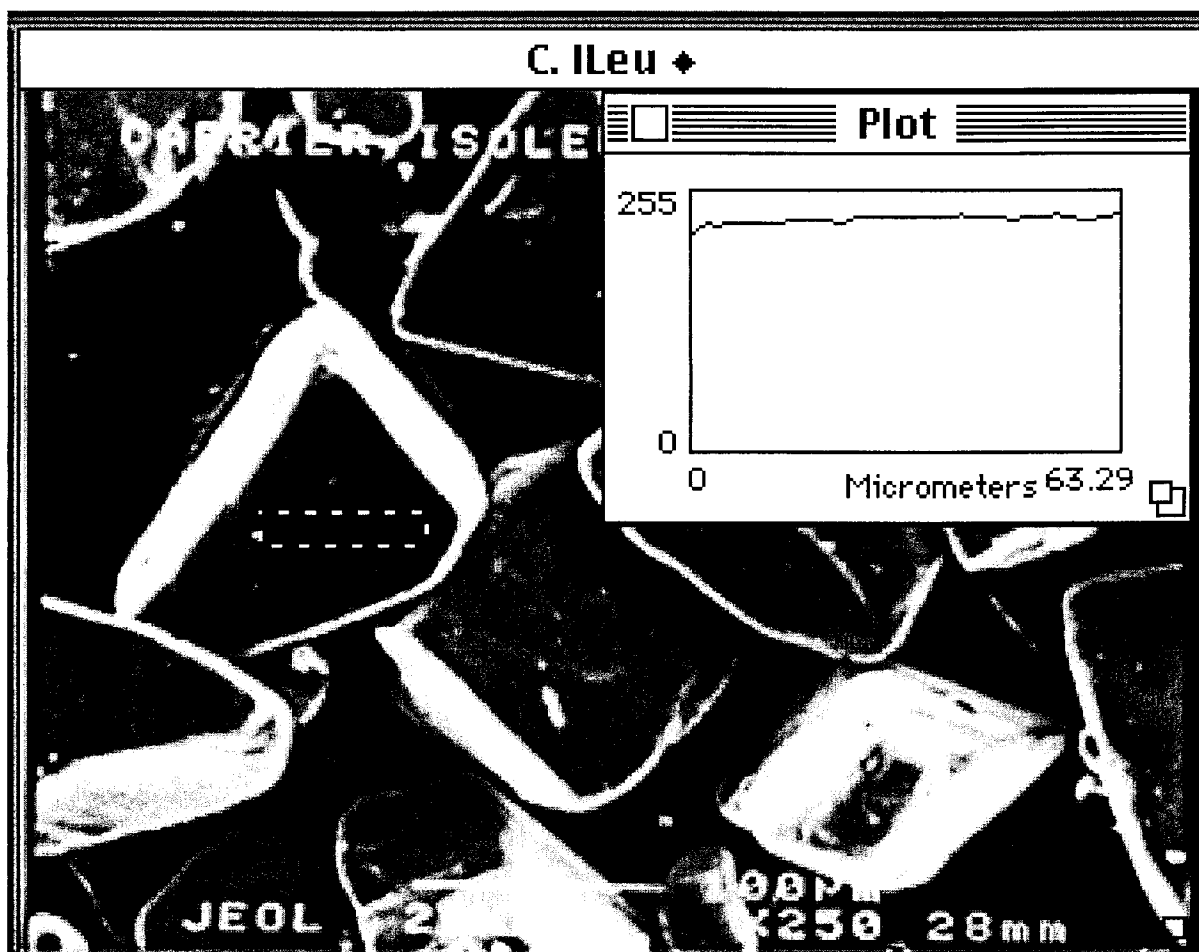


Figure 3

